

101033,055

(FILE 'HOME' ENTERED AT 09:40:34 ON 25 JUN 2004)

FILE 'REGISTRY' ENTERED AT 09:40:55 ON 25 JUN 2004

L1 1 S (169590-42-5)/RN
L2 1 S (76-42-6)/RN

FILE 'CAPLUS' ENTERED AT 09:41:40 ON 25 JUN 2004

L3 28 S L1 AND L2
L4 973 S L1
L5 732 S L2
L6 178 S L4 AND PAIN?
L7 8 S L6 AND SYNERG?
L8 56 S L4 AND (OPIOID? OR OPIATE? OR MORPHIN?)
L9 32 S L8 NOT L3
L10 8 S L1 AND PAIN? AND SYNERG?

FILE 'MEDLINE, SCISEARCH, BIOSIS' ENTERED AT 09:47:31 ON 25 JUN 2004

L11 2083 S L1
L12 2 S L11 AND PAIN? AND SYNERG?
L13 89 S L11 AND PAIN? AND (ACETAMINOPHEN? OR IBUPROFEN? OR ASPIRIN?)
L14 69 DUP REM L13 (20 DUPLICATES REMOVED)

FILE 'USPATFULL' ENTERED AT 09:50:44 ON 25 JUN 2004

L15 301 S L1
L16 221 S L15 AND (ASPIRIN? OR ACETAMIN? OR IBUPROFEN?)
L17 169 S L16 AND PAIN?
L18 58 S L17 AND (PAIN)/CLM
L19 23 S L18 AND SYNERG?

FILE 'USPATFULL' ENTERED AT 09:54:25 ON 25 JUN 2004

L20 21 S L1 AND L2
L21 1480 S (COX?)/CLM
L22 54 S (OPIOID? OR OPIAT? OR MORPHIN? OR OXYCOD?)/CLM AND L21

FILE 'WPIDS' ENTERED AT 09:57:47 ON 25 JUN 2004

L23 0 S L1

FILE 'CAPLUS' ENTERED AT 09:58:04 ON 25 JUN 2004

L24 973 S L1
L25 178 S L24 AND PAIN?

FILE 'STNGUIDE' ENTERED AT 09:59:42 ON 25 JUN 2004

FILE 'CAPLUS' ENTERED AT 10:01:37 ON 25 JUN 2004

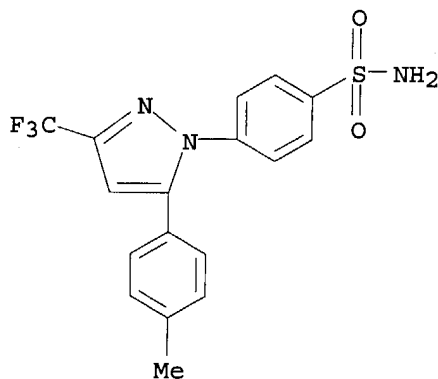
FILE 'STNGUIDE' ENTERED AT 10:03:17 ON 25 JUN 2004

FILE 'MEDLINE, SCISEARCH' ENTERED AT 10:04:01 ON 25 JUN 2004

L26 836 S L1
L27 155 S L26 AND PAIN?

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L1 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2004 ACS on STN
 RN 169590-42-5 REGISTRY
 CN Benzenesulfonamide, 4-[5-(4-methylphenyl)-3-(trifluoromethyl)-1H-pyrazol-1-yl]- (9CI) (CA INDEX NAME)
 OTHER NAMES:
 CN 4-[5-(4-Methylphenyl)-3-(trifluoromethyl)-1H-pyrazol-1-yl]benzenesulfonamide
 CN Celebrex
 CN Celecoxib
 CN Celocoxib
 CN SC 58635
 CN YM 177
 FS 3D CONCORD
 DR 184007-95-2, 194044-54-7
 MF C17 H14 F3 N3 O2 S
 CI COM
 SR US Adopted Names Council (USAN)
 LC STN Files: ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CANCERLIT, CAPLUS, CASREACT, CBNB, CEN, CHEMCATS, CIN, CSCHEM, DDFU, DIOGENES, DRUGU, EMBASE, HSDB*, IMSCOSEARCH, IMSDRUGNEWS, IMSPATENTS, IMSRESEARCH, IPA, MEDLINE, MRCK*, PHAR, PROMT, PROUSDDR, PS, RTECS*, SYNTHLINE, TOXCENTER, USAN, USPAT2, USPATFULL
 (*File contains numerically searchable property data)
 DT.CA Caplus document type: Book; Conference; Dissertation; Journal; Patent
 RL.P Roles from patents: ANST (Analytical study); BIOL (Biological study); PREP (Preparation); PROC (Process); PRP (Properties); RACT (Reactant or reagent); USES (Uses)
 RLD.P Roles for non-specific derivatives from patents: BIOL (Biological study); PREP (Preparation); PROC (Process); PRP (Properties); USES (Uses)
 RL.NP Roles from non-patents: ANST (Analytical study); BIOL (Biological study); MSC (Miscellaneous); PREP (Preparation); PROC (Process); PRP (Properties); RACT (Reactant or reagent); USES (Uses)
 RLD.NP Roles for non-specific derivatives from non-patents: BIOL (Biological study); PREP (Preparation); PROC (Process); PRP (Properties); USES (Uses)

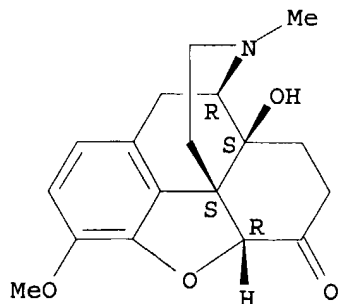


PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

962 REFERENCES IN FILE CA (1907 TO DATE)
 23 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
 973 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L2 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2004 ACS on STN
 RN 76-42-6 REGISTRY
 CN Morphinan-6-one, 4,5-epoxy-14-hydroxy-3-methoxy-17-methyl-, (5 α)-
 (9CI) (CA INDEX NAME)
 OTHER CA INDEX NAMES:
 CN Codeinone, 7,8-dihydro-14-hydroxy- (6CI, 7CI)
 CN Morphinan-6-one, 4,5 α -epoxy-14-hydroxy-3-methoxy-17-methyl- (8CI)
 OTHER NAMES:
 CN (-)-Oxycodone
 CN 14-Hydroxydihydrocodeinone
 CN 3-O-(Methyl)oxymorphone
 CN 6-Oxo-14-hydroxy-7,8-dihydrocodeine
 CN 7,8-Dihydro-14-hydroxycodeinone
 CN Dihydro-14-hydroxycodeinone
 CN Dihydrohydroxycodeinone
 CN Dihydrone
 CN NSC 19043
 CN Oxanest
 CN Oxicon
 CN Oxycodone
 CN Oxycodone
 CN Oxymorphone 3-methyl ether
 FS STEREOSEARCH
 MF C18 H21 N O4
 CI COM
 LC STN Files: ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, BEILSTEIN*,
 BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CANCERLIT, CAOLD, CAPLUS, CASREACT,
 CBNB, CEN, CHEMCATS, CHEMINFORMRX, CHEMLIST, CIN, CSCHEM, DDFU,
 DIOGENES, DRUGU, EMBASE, GMELIN*, HSDB*, IFICDB, IFIPAT, IFIUDB, IPA,
 MEDLINE, MRCK*, MSDS-OHS, NAPRALERT, NIOSHTIC, PHAR, PROMT, PROUSDDR,
 PS, RTECS*, SPECINFO, TOXCENTER, USAN, USPAT2, USPATFULL
 (*File contains numerically searchable property data)
 Other Sources: EINECS**, WHO
 (**Enter CHEMLIST File for up-to-date regulatory information)
 DT.CA Caplus document type: Conference; Journal; Patent
 RL.P Roles from patents: ANST (Analytical study); BIOL (Biological study);
 FORM (Formation, nonpreparative); MSC (Miscellaneous); PREP
 (Preparation); PROC (Process); PRP (Properties); RACT (Reactant or
 reagent); USES (Uses); NORL (No role in record)
 RLD.P Roles for non-specific derivatives from patents: BIOL (Biological
 study); PREP (Preparation); USES (Uses)
 RL.NP Roles from non-patents: ANST (Analytical study); BIOL (Biological
 study); FORM (Formation, nonpreparative); OCCU (Occurrence); PREP
 (Preparation); PROC (Process); PRP (Properties); RACT (Reactant or
 reagent); USES (Uses); NORL (No role in record)
 RLD.NP Roles for non-specific derivatives from non-patents: ANST (Analytical
 study); BIOL (Biological study)

Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

728 REFERENCES IN FILE CA (1907 TO DATE)
15 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
732 REFERENCES IN FILE CAPLUS (1907 TO DATE)
32 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

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L25 ANSWER 178 OF 178 CAPLUS COPYRIGHT 2004 ACS on STN

AN 1997:231026 CAPLUS

DN 126:264035

TI Synthesis and Biological Evaluation of the 1,5-Diarylpyrazole Class of Cyclooxygenase-2 Inhibitors: Identification of 4-[5-(4-Methylphenyl)-3-(trifluoromethyl)-1H-pyrazol-1-yl]benzenesulfonamide (SC-58635, Celecoxib)

AU Penning, Thomas D.; Talley, John J.; Bertenshaw, Stephen R.; Carter, Jeffery S.; Collins, Paul W.; Docter, Stephen; Graneto, Matthew J.; Lee, Len F.; Malecha, James W.; Miyashiro, Julie M.; Rogers, Roland S.; Rogier, D. J.; Yu, Stella S.; Anderson, Gary D.; Burton, Earl G.; Cogburn, J. Nita; Gregory, Susan A.; Koboldt, Carol M.; Perkins, William E.; Seibert, Karen; Veenhuizen, Amy W.; Zhang, Yan Y.; Isakson, Peter C.

CS Departments of Chemistry Inflammatory Diseases Research and Molecular Pharmacology, Searle Research and Development, Skokie, IL, 60077, USA

SO Journal of Medicinal Chemistry (1997), 40(9), 1347-1365

CODEN: JMCMAR; ISSN: 0022-2623

PB American Chemical Society

DT Journal

LA English

AB Sulfonamide-containing 1,5-diarylpyrazole derivs. were prepared and evaluated for their ability to block cyclooxygenase-2 (COX-2) in vitro and in vivo. Extensive structure-activity relationship work was carried out within this series, and a number of potent and selective inhibitors of COX-2 were identified. Since an early structural lead exhibited an unacceptably long plasma half-life, a number of pyrazole analogs containing potential metabolic sites were evaluated further in vivo in an effort to identify compds. with acceptable pharmacokinetic profiles. This work led to the identification of SC-58635 (celecoxib, I), which is currently in phase III clin. trials for the treatment of rheumatoid arthritis and osteoarthritis.

IT Pain

(hyperalgesia; diarylpyrazoles as cyclooxygenase 2 inhibitors)

IT 169590-42-5P 170569-50-3P 170569-69-4P 170569-75-2P
170569-83-2P 170569-85-4P 170569-88-7P 170569-91-2P 170570-25-9P
170570-80-6P 170570-81-7P 170571-00-3P 170571-05-8P 170571-29-6P
170571-71-8P 170571-92-3P 170571-97-8P 170572-00-6P 170572-05-1P
170572-08-4P 188816-97-9P 188816-98-0P 188816-99-1P 188817-00-7P
188817-07-4P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); RCT (Reactant); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); RACT (Reactant or reagent)
(diarylpyrazoles as cyclooxygenase 2 inhibitors)

L25 ANSWER 176 OF 178 CAPLUS COPYRIGHT 2004 ACS on STN
AN 1997:521520 CAPLUS
DN 127:171270
TI Outcome of specific COX-2 inhibition in rheumatoid arthritis
AU Lipsky, Peter E.; Isakson, Peter C.
CS Rheumatic Diseases Division, Department of Internal Medicine, University
Texas Southwestern Medical Center at Dallas, Dallas, TX, 75235-8884, USA
SO Journal of Rheumatology, Supplement (1997), 49(Progress toward a New Class
of Therapeutics: Selective COX-2 Inhibition), 9-14
CODEN: JRSUDX; ISSN: 0380-0903
PB Journal of Rheumatology
DT Journal
LA English

=> d 176 ab

L25 ANSWER 176 OF 178 CAPLUS COPYRIGHT 2004 ACS on STN
AB We reviewed data suggesting the hypothesis that specific inhibition of the
inducible isoform of cyclooxygenase, COX-2, would provide therapeutic
benefit in patients with rheumatoid arthritis (RA) with less
gastrointestinal toxicity and presented the results of a therapeutic trial
to test this hypothesis. Various doses of the selective COX-2 inhibitor,
celecoxib, or placebo were used to treat patients with RA in a 4 wk,
double blind, placebo controlled trial. Celecoxib provided significant
improvement in patient global assessment, morning stiffness, and the number
of **painful** and tender joints compared with placebo. In addition,
the number of withdrawals in celecoxib treated patients was significantly
less than in the placebo group. No significant adverse events and no
difference in the total number of adverse events were noted between the
placebo and celecoxib groups. At the doses employed, celecoxib inhibited
only COX-2 and not COX-1. Specific COX-2 inhibition with celecoxib causes
significant improvement in the signs and symptoms of RA.

L27 ANSWER 155 OF 155 MEDLINE on STN
AN 97393128 MEDLINE
DN PubMed ID: 9249645
TI Outcome of specific COX-2 inhibition in rheumatoid arthritis.
AU Lipsky P E; Isakson P C
CS Department of Internal Medicine, University of Texas Southwestern Medical
Center, Dallas 75235-8884, USA.
SO Journal of rheumatology, (1997 Jul) 24 Suppl 49 9-14. Ref: 21
Journal code: 7501984. ISSN: 0315-162X.
CY Canada
DT (CLINICAL TRIAL)
(CONTROLLED CLINICAL TRIAL)
Journal; Article; (JOURNAL ARTICLE)
General Review; (REVIEW)
(REVIEW, TUTORIAL)
LA English
FS Priority Journals
EM 199709
ED Entered STN: 19970916
Last Updated on STN: 20000303
Entered Medline: 19970904

L27 ANSWER 154 OF 155 MEDLINE on STN
AN 97393130 MEDLINE
DN PubMed ID: 9249647
TI **Pain** management in osteoarthritis: the role of COX-2 inhibitors.
AU Lane N E
CS Department of Medicine, University of California at San Francisco 94143,
USA.. nelane@itsa.ucsf.edu
NC AG05407 (NIA)
AR20684 (NIAMS)
SO Journal of rheumatology, (1997 Jul) 24 Suppl 49 20-4. Ref: 30
Journal code: 7501984. ISSN: 0315-162X.
CY Canada
DT Journal; Article; (JOURNAL ARTICLE)
General Review; (REVIEW)
(REVIEW, TUTORIAL)
LA English
FS Priority Journals
EM 199709
ED Entered STN: 19970916
Last Updated on STN: 20000303
Entered Medline: 19970904

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